



California Medical Device Recall Information



Recall Name

Baxter International Recalls INTRAVIA Containers Due to Particulate Matter

Recall Date	Product Description	Recalling Firm	Recall Reason
09/16/14	INTRAVIA Empty Containers with PVC Ports	Baxter Healthcare Corporation Deerfield, IL	<i>Due to complaints of particulate matter found inside the sterile fluid path.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	Two Lots: <ul style="list-style-type: none">INTRAVIA Container, 150 mL Capacity, Product Code: 2B8011 Lot: UR13D15112INTRAVIA Container, Empty 500 mL Capacity, Product Code: 2B8013 Lot: UR13K14095	CA , nationwide	<i>Lot # UR13D15112:</i> distributed between April 26 and June 20, 2013; <i>Lot # UR13K14095:</i> distributed between November 27, 2013 and March 10, 2014.

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm423933.htm>